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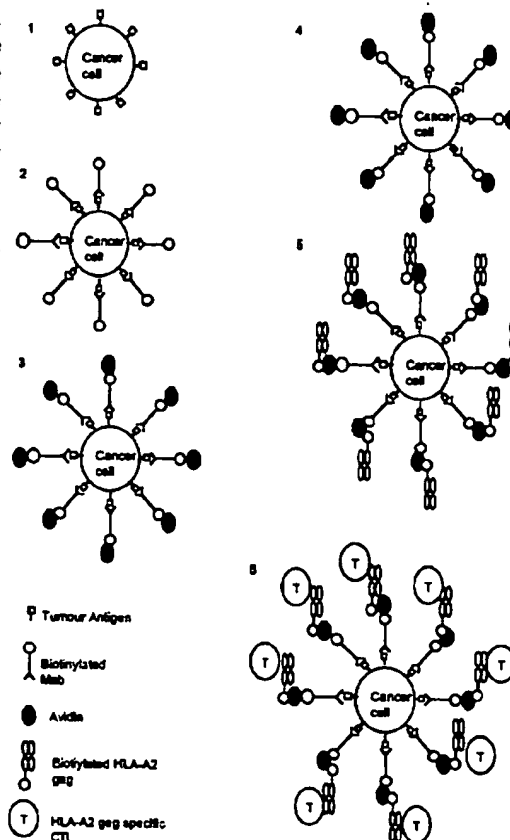
(58) Field of Search

ONLINE: WPI, EPODOC, PAJ, BIOSIS, CAPLUS,  
BIOTECHABS, CANCERLIT, EMBASE, LIFESCI,  
MEDLINE

(54) Abstract Title

**Chimeric protein complexes comprising HLA class I antigens**

(57) There are described HLA class I molecules linked to means to selectively attach the molecule to the cell. Attachment means are preferably antibodies and more preferably antibodies against tumour antigens to facilitate the deposition of class I molecules on tumour cells. The class I molecule and cell attachment means are preferably linked via an avidin/biotin bridge. The class I molecule may be allogeneic. Alternatively autologous HLA molecules may be used to increase Ag presentation by tumour cells, expressing low amounts of class I Ag, or antigen presenting cells (APC's). The HLA molecule has intact T cell binding portions.



At least one drawing originally filed was informal and the print reproduced here is taken from a later filed formal copy.

The claims were filed later than the filing date but within the period prescribed by Rule 25(1) of the Patents Rules 1995.

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Tumour Antigen

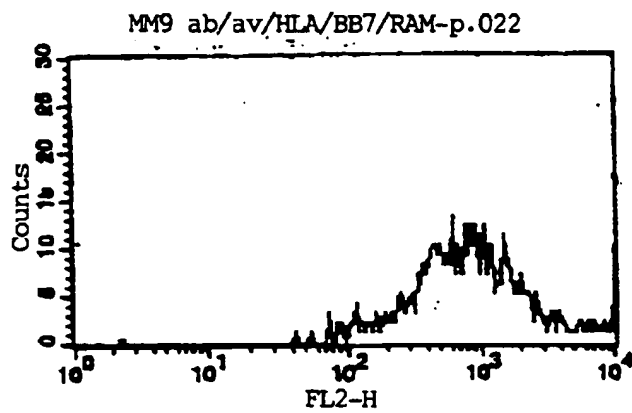
Biotinylated Mab

Avidin

Biotinylated HLA-A2 gag

HLA-A2 gag specific CTL

2/9.



antibody/avidin/HLA/  
BB7.2/RAH-PE

MM9 melanoma 12.97

3/9.

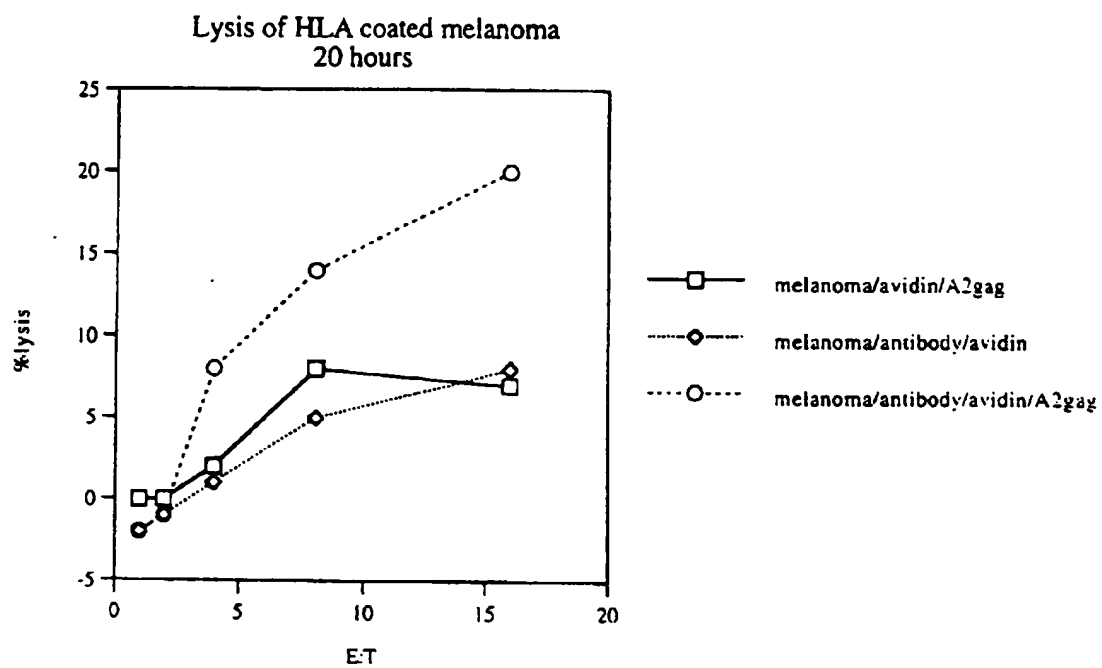
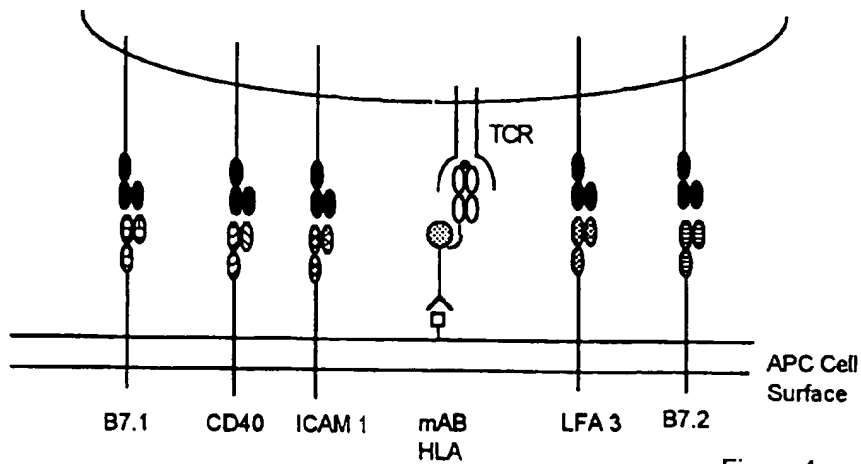
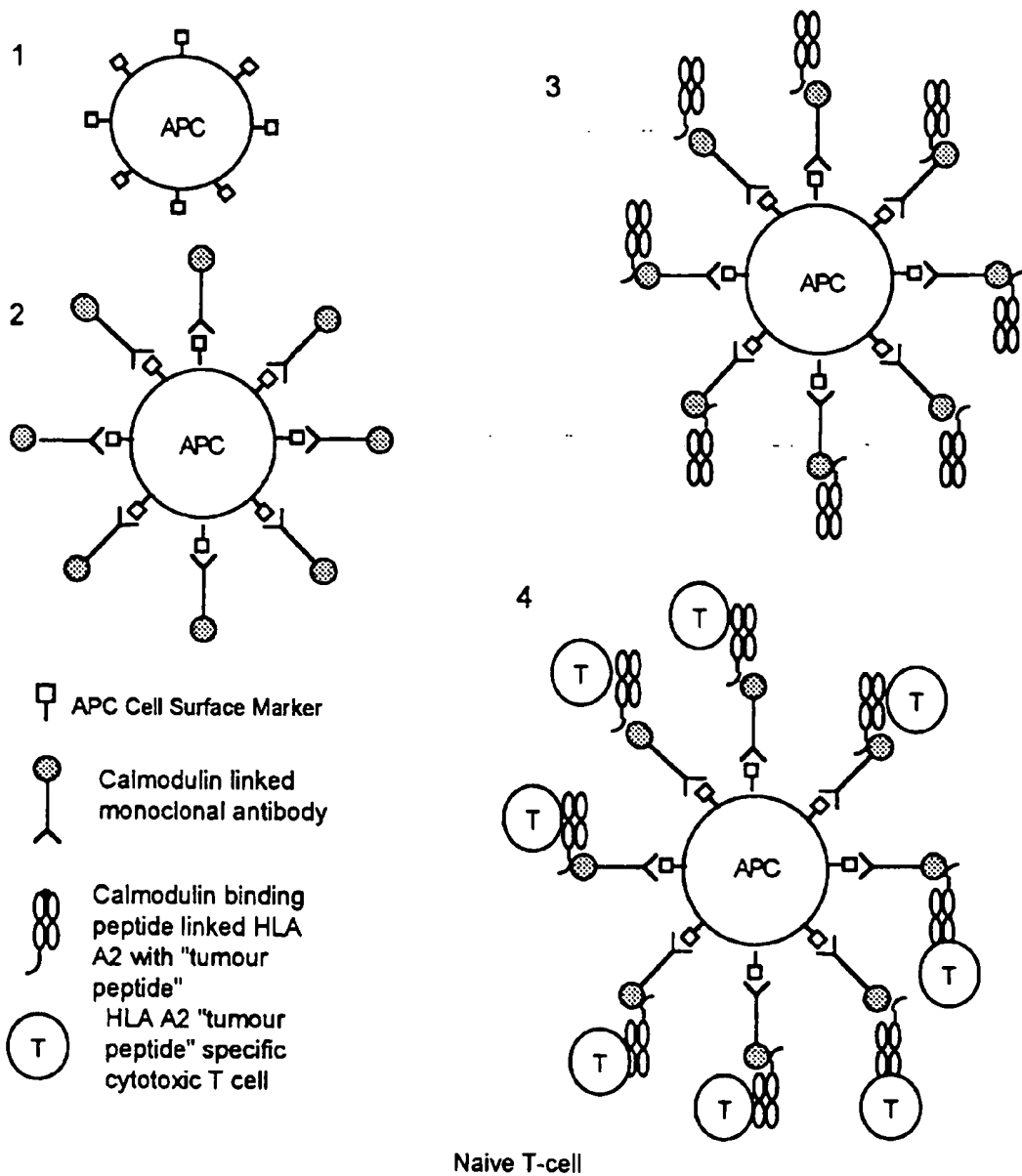


Figure 3

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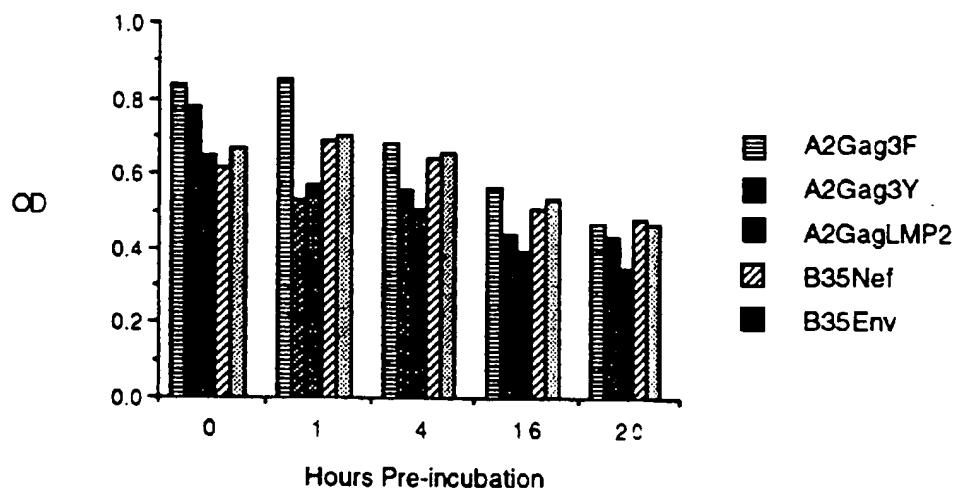


FIGURE 5

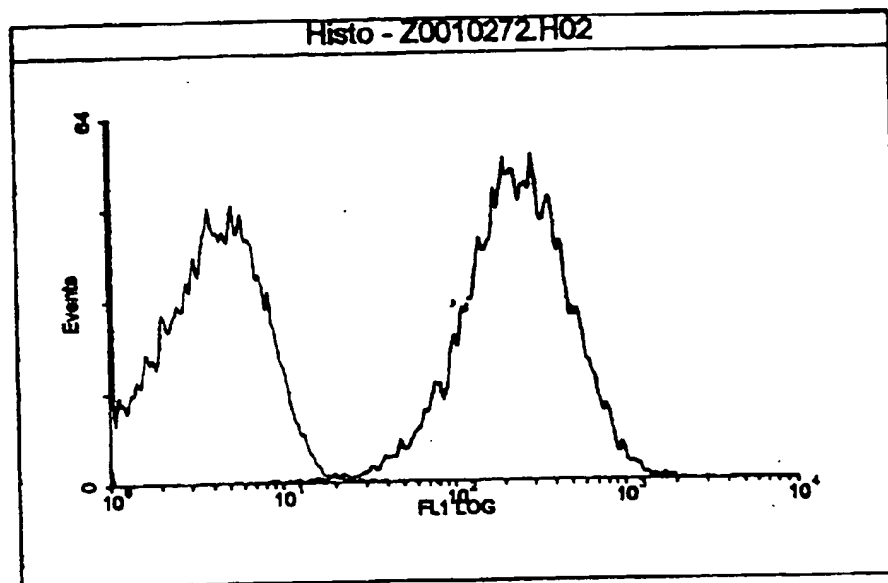


Figure 6

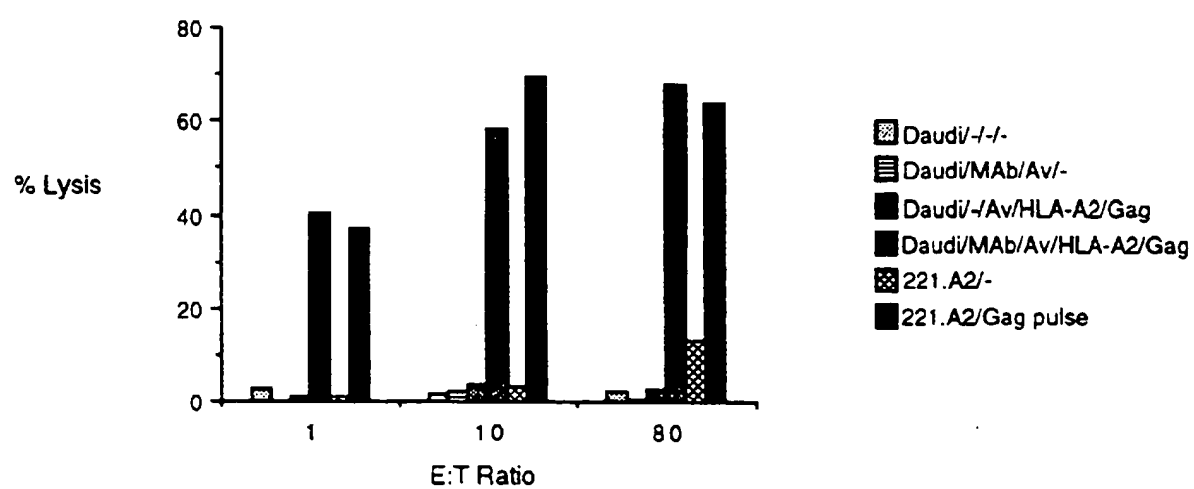


FIGURE 7



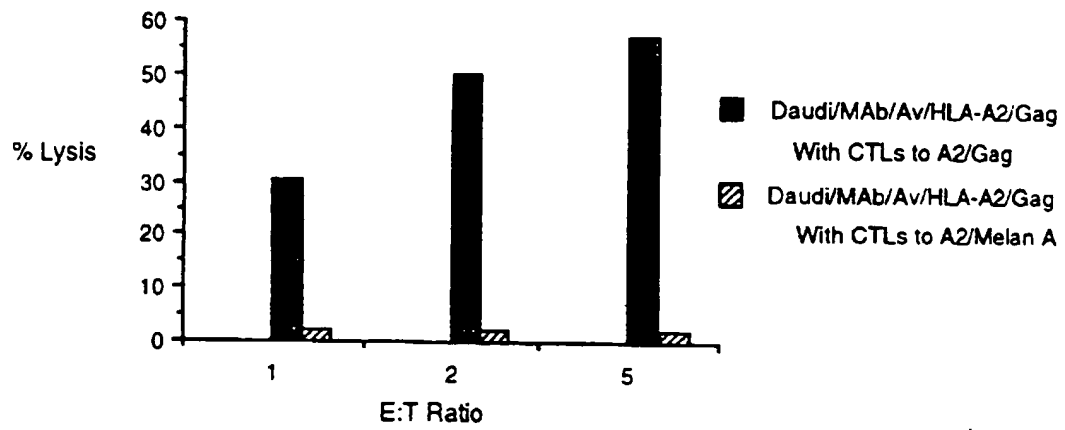


FIGURE 8

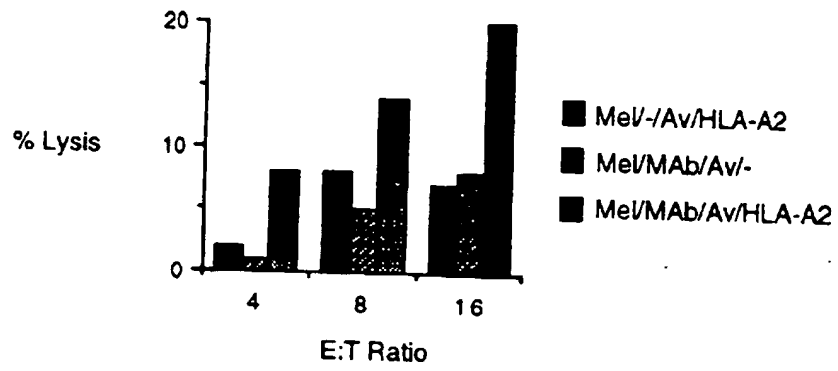


FIGURE 9

NOVEL METHOD II

This application relates to means for producing or enhancing an immunological  
5 response against a target cell, through the attachment of an immunogenic HLA  
class I molecule thereto. The invention is of potential use in the prevention and  
treatment of malignant diseases including cancer and leukaemia, infectious  
diseases including viral infections such as HIV, bacterial infections including  
tuberculosis, and parasitic infections including malaria.

10

Cytotoxic T cells in the cellular immune system are responsible for recognising  
cells that display "foreign" markings, and triggering an immunological response  
against such cells. Each cytotoxic T cell expresses a number of cell surface  
recognition receptors, which recognition receptors all possess precise specificity  
15 for a particular "foreign" peptide sequence, which recognition receptors are  
adapted to bind to HLA class I molecules expressed on the surface of cells  
scanned by the T cell. HLA class I molecules are cell surface molecules which  
possess a peptide binding groove exposed on the external surface of the cell,  
which groove is arranged under normal circumstances to bind a peptide derived  
20 from the interior of the cell. When a recognition receptor on a cytotoxic T cell  
binds to an HLA class I molecule on the surface of a scanned cell, the

recognition receptor is enabled to contact the peptide binding groove of the HLA class I molecule and interact with any peptide contained therein. If this peptide matches the specificity of the recognition receptor, the T cell is said to recognise the scanned cell, and may consequently trigger an immunological  
5 response against said scanned cell.

Cytotoxic T cells of various specificities within a host immune system are also able to recognise and trigger an immunological response against a cell exhibiting an HLA class I molecule which is of a different allotype from the  
10 HLA class I molecules of the host cells. An immunological response of this kind is known as an "alloreactive" response.

An immunological response against a cell usually results in the lysis of the cell and/or the local release of cytokines. It has however been observed that  
15 cytotoxic T cells do not trigger the lysis of so-called antigen presenting cells (APCs) in this way. Instead, the immunological response triggered by T cell recognition of an HLA class I molecule on the surface of an antigen presenting cell results in the direct selective proliferation of the cytotoxic T cell. The host immune system consequently becomes immunised against any cells exhibiting  
20 the foreign peptide recognised by the surface recognition receptors on this T cell.

It is recognised that the effector mechanisms of the cellular immune system could be a powerful tool in the prevention and treatment of many illnesses, including malignant processes and infectious and auto-immune diseases, including cancer. A small number of the HLA class I molecules on a tumour cell surface may be found to bind peptides which are selectively expressed or over-expressed in tumour cells and are capable of being recognised by cytotoxic T cells in the immune system. Such peptides may furthermore be tumour specific, being found only infrequently, or not at all, on the HLA class I molecules of non-tumour cells. An example of one such tumour specific peptide is the HMW-MAA antigen found on melanoma cells. However, the number of HLA molecules presenting such peptides is generally too small to stimulate an effective immunological response against the tumour cell. Moreover, such peptides are rarely, if ever, presented by HLA class I molecules on the surface of APCs.

15

Attempts to enhance the response of the cellular immune system to tumour cells have hitherto focused on increasing tumour cell immunogenicity. In particular, various efforts have been made to produce high-level expression of immunogenic HLA class I molecules on the surface of tumour cells, through the techniques of gene therapy. The delivery of cDNA encoding an HLA class I gene containing an immunogenic peptide in the leader sequence of the HLA

20

molecule has been described in Kang (Cancer Res. 57, 1997, 202-205).

Meanwhile, Stopeck (J Clinical Oncology 15, 1997, 341-349) describes the transfection of allogeneic HLA class I in patients with melanoma. This work has demonstrated some response in clinical trials, but has also highlighted the difficulties involved in targeting tumour cells at multiple sites in vivo through the techniques of gene therapy.

The present application sets out to provide improved means for producing or enhancing an immunological response against a target cell, and to provide an improved method for treating or preventing cancer and other malignant, infectious or auto-immune diseases.

Accordingly, in one aspect of the present invention there is provided a complex comprising an HLA class I molecule or fragment thereof, which HLA class I molecule or fragment thereof comprises a T cell binding portion, and attaching means for selectively attaching said HLA class I molecule or fragment thereof to a target cell.

In another aspect of the present invention there is provided a method of attaching an HLA class I molecule or fragment thereof to a target cell, which HLA class I molecule or fragment thereof comprises a T cell binding portion,

comprising the step of introducing to said target cell said HLA class I molecule or fragment thereof and attaching means for selectively attaching said HLA class I molecule or fragment thereof to the target cell.

- 5 In yet another aspect of the present invention, there is provided a pharmaceutical composition comprising an HLA class I molecule or fragment thereof, which HLA class I molecule or fragment thereof comprises a T cell binding portion; attaching means for selectively attaching said HLA class I molecule or fragment thereof to a target cell; and an appropriate excipient or  
10 carrier.

The HLA class I molecule or fragment thereof may bind a peptide, which peptide is arranged to be presented for T cell recognition by said HLA class I molecule or fragment thereof. Said peptide may be attached to the HLA class I  
15 molecule or fragment thereof in accordance with the method described in Garboczi (PNAS 89, 1992, 3429-3433).

The attaching means preferably comprises a linking polypeptide with high specific affinity for a target cell specific molecule on the surface of the target  
20 cell. By "target cell specific molecule" herein is meant any molecule that is characteristically expressed or over-expressed on the surface of the target cell.

By way of example, in cancer cells said "target cell specific molecule" could include any of the following tumour associated antigens: carcinoembryonic antigen, placental alkaline phosphatase, polymorphic epithelial mucin, human chorionic gonadotrophin, CD20, prostate specific antigen, ca-125, HMW-MAA  
 5 and others.

Conveniently, the linking polypeptide will comprise an antibody, preferably a monoclonal antibody, raised against said target cell specific molecule (Riethmuller and Johnson, Curr. Opin. Immunol. 4, 1992, 647-655). Suitable  
 10 antibodies for this purpose include C46, 85A12, H17E2, HMFG1, W14, 1F5, 225.28s (Buraggi 1985 Cancer Res. 45, 3378-3387), and others. Deposits of the immortalised hybrids producing these antibodies have been made at the American Type Culture Collection, Rockville MD, USA. Further examples of antibodies are described in Maloney et al (Blood 84, 1994, 2457-2466),  
 15 Riethmuller et al (Lancet 343, 1994, 1177-1183) and Hird et al (Br. J. Cancer 68, 1993, 403-406).

Said linking polypeptide may comprise an antibody raised against a target cell specific molecule and a coupling system for coupling said antibody to said  
 20 HLA class I molecule or fragment thereof. The coupling system may comprise a two- or three-step chain of well-characterised paired small molecules, joined



to the antibody and the HLA class I molecule so as to form a stable bridge between the two. Examples of paired small molecules which might be used in this connection include (but are not limited to) biotin and avidin/streptavidin (Moro, 1997 Cancer Res. 57, 1922-1928; Altman et al, Science 274, 1996, 94-  
5 96), and calmodulin and calmodulin binding peptides (Neri, 1996, J. Invest. Dermatol. 107, 164-170). Alternatively, said linking polypeptide may comprise an antibody raised against a target cell specific molecule, which antibody is adapted to be attached directly to said HLA class I molecule or fragment thereof.

10

In a further possible embodiment of the invention, said complex may comprise a recombinant protein, which recombinant protein includes a moiety comprising said HLA class I molecule or fragment thereof, and a moiety comprising said attaching means.

15

The HLA class I molecule or fragment thereof may be purified from plasma or platelets or made recombinantly. The HLA class I molecule or fragment thereof may further be arranged to bind and present for T cell recognition a defined peptide of choice, such as a viral, bacterial, parasitic, or tumour-specific

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peptide. Attachment of the HLA class I molecule or fragment thereof to the target cell may be achieved by introducing said HLA class I molecule or

fragment thereof and said attaching means to the vicinity of the target cell. The target cell may be a culture cell in vitro, but will advantageously be in the body of a patient. Preferably, the target cell will be arranged to be contacted by a cytotoxic T cell, which cytotoxic T cell is adapted to recognise said HLA class I molecule or fragment thereof either as being of a mismatched allotype or as binding a foreign peptide, and which cytotoxic T cell is capable of triggering an immunological response against said target cell.

In one embodiment of the present invention the target cell is of a type which may be lysed as a result of an immunological response thereagainst. Advantageously, the target cell is a tumour cell or any diseased or foreign cell the presence of which is undesired in a patient, such as a cancer cell, leukaemia cell, a cell infected with the HIV virus or with any other microbe or virus, a cell responsible for detrimental activity in auto-immune disease, and so on. In order to accelerate the triggering of an immunological response against said target cell in a patient, said HLA class I molecule or fragment thereof will preferably be capable of producing a powerful immune response from the cellular immune system of the patient. Accordingly, said HLA class I molecule or fragment thereof may bind a viral or microbial peptide, preferably a viral or microbial peptide to which the patient is likely to have had previous exposure. In particular, said HLA class I molecule or fragment thereof may bind an

influenza virus peptide, a measles virus peptide, an Epstein-Barr virus peptide, in particular an Epstein-Barr virus peptide comprising the RAKFFQLL epitope of the lytic protein BZLF1, a Cytomegalovirus peptide, or a tetanus toxoid peptide. Alternatively, said HLA class I molecule or fragment thereof may bind  
5 any peptide which already has a strong cytotoxic T cell response or which is capable of inducing a powerful immune response. The allotype of said HLA class I molecule or fragment thereof may additionally or alternatively be different from the allotype of the HLA class I molecules of the patient, so that an alloreactive response may additionally or alternatively be triggered against  
10 said target cell.

In another embodiment of the invention the target cell is an antigen presenting cell (APC). Recognition by a cytotoxic T cell of an HLA class I molecule or fragment thereof attached to said APC may result in direct and selective  
15 proliferation of the cytotoxic T cell. Accordingly, said HLA class I molecule or fragment thereof will advantageously be adapted to present for T cell recognition a tumour specific peptide as defined above, or a viral peptide, or a bacterial peptide, or a parasitic peptide, or any peptide which is exclusively or characteristically presented by HLA class I molecules on the surface of  
20 diseased, malignant or foreign cell; the presence of which is undesirable in a patient. Peptides linked to malignant conditions have been characterised

(Brossart, 1998 Cancer Res. 58, 732-736 and Lucas, 1998 Cancer Res. 58, 743-752), as have peptides of parasitic origin (Khusmith, 1991 Science 252, 715-718) The attachment of an HLA class I molecule or fragment thereof to an APC, in accordance with the present invention, may be used for in vivo  
5 immunisation against cells presenting a given peptide, or ex vivo production of cytotoxic T cells of a particular specificity.

Where the target cell is a tumour cell or microbially infected cell, the pharmaceutical composition of the present invention may be used for the  
10 treatment of a tumour or microbial disease respectively, and there is provided a method of treating a tumour or microbial disease in a patient, comprising the step of administering to a patient in need thereof an effective amount of said pharmaceutical composition.

15 It must be noted that whilst many tumour types express tumour associated antigens, heterogeneity in the level of expression does occur, so some tumour cells may not be targeted by antibody and lysed directly. However, in vitro data from the analogous antibody-superantigen system shows that the high local levels of cytokines released by activated T cells can lead to the death of  
20 untargeted bystander tumour cells (Dohlsten et al, Int. J. Cancer 54, 1993, 482-488). It is likely that similar effects will occur in a targeting system using

MHC class I/peptide complexes. Similarly, it is possible that the presence of activated cytotoxic T cells releasing cytokines in the tumour may lead to enhancement of a specific anti-tumour immune response.

Where the target cell is an APC and the HLA class I molecule or fragment thereof binds a tumour-specific peptide or any peptide which is exclusively or characteristically presented by HLA class I molecules on the surface of a virally, bacterially, parasitically or microbially infected cell, the pharmaceutical composition of the present invention may be used for immunising against the tumour or viral, bacterial, parasitic or microbial infection respectively, and there is provided a method of immunising against a tumour or viral, bacterial, parasitic or microbial infection in a patient, comprising the step of administering to a patient in need thereof an effective amount of said pharmaceutical composition.

The response of said patient may be improved by in vivo cytokine support, or by the infusion of antigen-specific cytotoxic T cells expanded ex vivo.

Transient immunosuppression (Ledermann et al, Int. J. Cancer 47, 1991, 659-664) may be used to minimise the immunogenic response of a patient to components of the targeting system such as the avidin bridge.

20

The administration of said pharmaceutical composition may be by way of oral,

sublingual, transdermal or parenteral administration.

Said effective amount of the pharmaceutical composition will depend on factors such as the nature and severity of the disorder being treated and on the weight,  
5 age and condition of the patient.

For oral or parenteral administration, it is greatly preferred that the pharmaceutical composition is administered in the form of a unit-dose composition, such as a unit dose oral or parenteral composition.

10

Such compositions are prepared by admixture and are suitably adapted for oral or parenteral administration, and as such may be in the form of tablets, capsules, oral preparations, powders, granules, lozenges, reconstitutable powders, injectable and liquid infusible solutions or suspensions or  
15 suppositories.

20

Tablets and capsules for oral administration are usually presented in a unit dose, and contain conventional excipients such as binding agents, fillers, diluents, tableting agents, lubricants, disintegrants, colourants, flavourings, and wetting agents. The tablets may be coated according to well known methods in the art.

Suitable fillers for use include cellulose, mannitol, lactose and other similar agents. Suitable disintegrants include starch, polyvinylpyrrolidone and starch derivatives such as sodium starch glycolate. Suitable lubricants include, for example, magnesium stearate. Suitable pharmaceutically acceptable wetting agents include sodium lauryl sulphate.

These solid oral compositions may be prepared by conventional methods of blending, filling or tableting. Repeated blending operations may be used to distribute the active agent throughout those compositions employing large quantities of fillers. Such operations are, of course, conventional in the art.

Oral liquid preparations may be in the form of, for example, aqueous or oily suspensions, solutions, emulsions, syrups, or elixirs, or may be presented as a dry product for reconstitution with water or other suitable vehicle before use.

Such liquid preparations may contain conventional additives such as suspending agents, for example sorbitol, syrup, methyl cellulose, gelatin, hydroxyethylcellulose, carboxymethyl cellulose, aluminium stearate gel or hydrogenated edible fats, emulsifying agents, for example lecithin, sorbitan monooleate, or acacia; non-aqueous vehicles (which may include edible oils), for example, almond oil, fractionated coconut oil, oily esters such as esters of glycerine, propylene glycol, or ethyl alcohol; preservatives, for example methyl

or propyl p-hydroxybenzoate or sorbic acid, and if desired conventional flavouring or colouring agents.

Oral formulations also include conventional sustained release formulations,  
5 such as tablets or granules having an enteric coating.

For parenteral administration, fluid unit dose forms may be prepared comprising a sterile vehicle. The components of the composition, depending on the vehicle and the concentration, can be either suspended or dissolved.  
10 Parenteral solutions are normally prepared by dissolving the components of the composition in a vehicle and filter sterilising before filling into a suitable vial or ampoule and sealing. Advantageously, adjuvants such as a local anaesthetic, preservatives and buffering agents are also dissolved in the vehicle. To enhance the stability, the composition may be frozen after filling into the vial  
15 and the water removed under vacuum.

Parenteral suspensions are prepared in substantially the same manner except that the compound may be suspended in the vehicle instead of being dissolved and sterilised by exposure to ethylene oxide before suspending in the sterile  
20 vehicle. Advantageously, a surfactant or wetting agent may be included in the composition to facilitate uniform distribution of the compound of the invention.



As is common practice, the compositions will usually be accompanied by written or printed directions for use in the treatment concerned.

Following is a description, by way of example only, and with reference to the accompanying drawings, of methods of putting the present invention into effect.

In the drawings :-

Figure 1 shows a diagram showing the method/idea for delivering HLA molecules to the surface of tumour cells.

Figure 2 shows a FACs analysis of HLA-A2-ve Mel 2 melanoma cells treated with biotin-conjugated monoclonal antibody 225.28s, avidin, biotin-conjugated HLA-A2/gag complexes, anti-HLA-A2 monoclonal antibody BB7.2 and phycoerythrin-conjugated rabbit anti-mouse antibody.

Figure 3 shows the results of a T cell cytotoxicity chromium release assay with Mel 1 cells treated with the delivery system of biotin-conjugated monoclonal antibody 225.28s, avidin, and biotin-conjugated HLA-A2/gag complexes. These cells were incubated with HLA-A2/gag specific cytotoxic T cells with effector/target ratios of 0:1 - 20:1 for 20 hours.

Figure 4 shows a diagram showing the method/idea for delivering HLA class I/peptide complexes to antigen presenting cells.

Figure 5 shows the results of an ELISA assay, described in Example 2 below, for demonstrating the stability of various MHC class I/peptide complexes at 37°C. The results shown are the mean of assays performed on each sample in triplicate.

Figure 6 shows a FACS analysis of HLA-class I deficient Daudi cells targeted with HLA-A2 via biotinylated anti-CD20 mAb. Trace 1 (lefthand trace) corresponds to native untargeted Daudi cells. Trace 2 (righthand trace) corresponds to Daudi cells targeted with mAb/avidin/HLA-A2/gag/FITC anti-MHC class I. Mean fluorescence trace 1 = 0.31, mean fluorescence trace 2 = 24.3 (arbitrary fluorescence units).

Figure 7 shows the results of a four hour chromium release assay, described in Example 2 below, in which HLA class I-deficient Daudi cells targeted with various components of the HLA-A2/gag delivery system were incubated with HLA-A2/gag specific cytotoxic T cell clones. A comparison was made with native and peptide-pulsed 221.A2 cells (HLA-A2 +ve).

Figure 8 shows the results of a four hour chromium release assay, described in Example 2 below, in which HLA-A2/gag targeted Daudi cells were incubated with HLA-A2/gag-specific and HLA-A2/Melan A-specific cytotoxic T cell clones.

5

Figure 9 shows the results of a twenty hour chromium release assay, described in Example 2 below, in which HLA-A2 +ve SK29.Mel cells were incubated with HLA-A2/gag specific cytotoxic T cell clones.

#### 10 Example 1

The following components were used:

Target cells :

15 A human melanoma cell line Mel 1, deposited at the Department of Immunology, Institute of Molecular Medicine, Oxford, that carries the HLA class I allotype HLA-A2. The cell line was grown in standard RPMI tissue culture media.

20 A human melanoma cell line Mel 2, deposited at the Department of Immunology, Institute of Molecular Medicine, Oxford, that does not carry the HLA class

I allotype HLA-A2. The cell line was grown in standard RPMI tissue culture media.

Attaching means : A monoclonal antibody 225.28s (Buraggi 1985 Cancer Res.  
 5 45, 3378-3387) that binds to the HMW-MAA antigen on human melanoma cells. Biotin is chemically conjugated onto this antibody as described in Bayer 1990, Methods Embryology 184, 138-160.

Pure hen egg avidin obtained commercially from  
 10 Societa Prodotti Antibiotici, Milan, Italy.

HLA : Biotin conjugated recombinant HLA class I allotype HLA-A2 molecules, as described in Altman 1996, Science 274, 94-96, further containing the "gag" peptide that is part of the HIV virus. This peptide  
 15 comprises the amino acid sequence -SLYNTVATL-. Methods for the preparation/isolation thereof are described in Johnson 1991, J Immunol 147, 1512. The "gag" peptide was attached to the HLA-A2 molecules as described in Garboczi 1992, PNAS 89, 3429-3433.

20 T cells : HLA-A2/gag specific cytotoxic T cells obtained from an A2+ve HIV patient as described in Altman 1994,

In order to establish the ability of the attaching means to cause display of the HLA class I molecules on the surface of Mel 2 target cells, approximately  
5 200,000 cells were first incubated with biotin conjugated monoclonal antibody 225.28s at a final concentration of 20µg/ml at 37°C for 30 minutes. Following this the cells were washed in tissue culture media (RPMI 1640, obtainable from Gibco, Scotland). The Mel 2 cells were then incubated with avidin at a final concentration of 10µg/ml for 10 minutes at 37°C and washed in tissue culture  
10 media. Finally, the Mel 2 cells were incubated with biotin conjugated HLA class I HLA-A2/gag molecules at a final concentration of 20µg/ml at 37°C for 20 minutes.

The binding of recombinant HLA-A2 to the treated Mel 2 cells was shown by  
15 the attachment of anti-HLA-A2 monoclonal antibody BB7.2 (Santos-Aguado 1988, J. Immunol 141, 2811-2818) following incubation with BB7.2 antibody at a final concentration of 10µg/ml at 37°C for 30 minutes. After washing in tissue culture media the cells were incubated with phycoerythrin conjugated rabbit anti-mouse antibody (Sigma, Poole, UK) at a final concentration of  
20 10µg/ml for 30 minutes at 37°C and analysed in a Becton Dickson Facscan machine. The result of this analysis is shown in Fig 2 which demonstrates a

positive signal indicating the presence of HLA-A2 molecules attached to the surface of the Mel 2 cells.

A chromium release T cell cytotoxicity assay was then performed in order to establish the ability of HLA-A2/gag specific T cell clones to lyse Mel 1 cells coated with HLA-A2/gag in accordance with the present method.

Approximately  $10^6$  Mel 1 cells were first pre-incubated with  $1.85\mu\text{Bq}$   $\text{Na}_2^{51}\text{CrO}_4$  (obtained from Amersham International, Amersham, UK) for 1 hour at  $37^\circ\text{C}$ . The pre-incubated Mel 1 cells were then incubated with biotin conjugated monoclonal antibody 225.28s at a final concentration of  $20\mu\text{g/ml}$  at  $37^\circ\text{C}$  for 30 minutes, and washed in tissue culture media. Following this, the Mel 1 cells were incubated with avidin at a final concentration of  $10\mu\text{g/ml}$  for 10 minutes at  $37^\circ\text{C}$  and then washed again in tissue culture media. The Mel 1 cells were then incubated with biotin conjugated HLA class I HLA-A2/gag molecules at a final concentration of  $20\mu\text{g/ml}$  at  $37^\circ\text{C}$  for 20 minutes and washed with tissue culture media.

Having been coated with HLA class I HLA-A2/gag, the chromium-treated Mel 1 cells were then incubated with HLA-A2/gag specific cytotoxic T cells in ratios of 0:1 to 20:1 of effector to target cells at  $37^\circ\text{C}$  for 20 hours. Lysis of Mel 1 cells treated with  $\text{Na}_2^{51}\text{CrO}_4$  results in the release of radioactive

chromium, which may be detected by analysis in a scintillation counter.

In order to establish the percentage of Mel 1 cells lysed following incubation with HLA-A2/gag specific cytotoxic T cells, the following measurements were taken: background release of chromium from the Mel 1 cells in media alone ("M"); release of chromium from the Mel 1 cells following incubation with the T cells ("E"); release of chromium from the Mel 1 cells following final treatment with 5% Triton X-100 detergent ("T"). Treatment with detergent will cause the lysis of all the remaining intact Mel 1 cells.

% Mel 1 lysis by cytotoxic T cells was calculated as follows:

$$\% \text{ lysis} = 100 \times \frac{(E - M)}{(T - M)}$$

This analysis was carried out on Mel 1 cells treated with biotin-conjugated 225.28s, avidin, and biotin-conjugated HLA-A2/gag. As a control, the analysis was also carried out on Mel 1 cells treated with biotin-conjugated 225.28s and avidin alone, and on Mel 1 cells treated with avidin and biotin-conjugated HLA-A2/gag alone.

The key results of this analysis are illustrated in Figure 3, which indicates that

significant lysis (20%) of Mel 1 cells by HLA-A2/gag specific cytotoxic T cells occurs only when the Mel 1 cells have been treated with all the components of the attaching and delivery means of the present invention (ie biotin-conjugated 225.28s monoclonal antibodies, avidin, and biotin-conjugated HLA-A2/gag). No  
5 significant increase in cell lysis over background levels was observed in either of the control runs.

### Example 2

10 The following components were used:

Target cells :	The Daudi B cell line (MHC class I-negative) melanoma line SK-mel-29 (HLA-A2.1-positive), .221/A2 (HLA-A2.1-positive), were maintained in 15 RPMI media with 10% fetal calf serum and antibiotics in a 37°C incubator with 5% CO <sub>2</sub> .
Attaching means :	Monoclonal antibodies 225.28s (Buraggi 1985 <u>Cancer Res.</u> 45, 3378-3387) and 2H7 that bind to the 20 HMW-MAA antigen. Biotin is chemically conjugated onto these antibodies as described in Bayer 1990,



Methods Embryology 184, 138-160.

Pure hen egg avidin obtained commercially from  
Societa Prodotti Antibiotici, Milan, Italy.

5

HLA :

Biotinylated complexes of recombinant MHC class I  
and peptide were produced as described previously  
(Altman et al, Science 274, 1996, 94-96; Ogg et al,  
Science 279, 1998, 2103). Prokaryotic expression of  
B2M and MHC class I heavy chain, modified by the  
C terminal addition of a target sequence for the  
biotin ligase enzyme BirA, was followed by  
inclusion body purification. Following refolding of  
heavy chain and B2M around specific peptide,  
complexes of 45kD were isolated by gel filtration,  
biotinylated overnight using BirA in the presence of  
ATP, Mg<sup>2+</sup> and biotin, and then purified by gel  
filtration and anion exchange.

10

15

20 T cells :

Human cytotoxic T cell clones 010 (specific for  
HLA-A2/gag 77-85 = SLYNTVATL (Parker et al, *J*

Immunol. 149, 1992, 3580-3587)) and IF9 (specific for HLA-A2/melan-A 26-35 = EAAGIGILTV (Romero et al, J. Immunol. 159, 1997, 2366) were maintained in media supplemented with 5% human serum and IL-2 100 IU/ml.

5

The stability of the MHC class I/peptide complexes was first established by an ELISA assay. Various MHC class I/peptide complexes, including HLA-A2/Gag3Y, HLA-A2/Gag3F, HLAA2/Lmp2, HLA-B35/Env and HLA-B35/nef, were prepared as outlined above, and were pre-incubated at 10ug/ml in tissue culture media for 0-20 hours at 37°C. ELISA plates were coated with the mAb W6/32 (5ug/ml in carbonate buffer pH 9.6 overnight at 4°C) which recognises conformationally correct MHC class I molecules (Parham, 1979), and then blocked by incubation in 1% bovine serum albumin for 2 hours at 37°C. The MHC class I/peptide complexes were incubated for 30 minutes with the ELISA plates at room temperature, and binding was detected with rabbit anti-human B2 microglobulin followed by alkaline phosphatase conjugated goat anti-rabbit immunoglobulin, and substrate. All incubations were separated by extensive washes in PBS.

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Absorbances at 600nm were measured in a Titertek Multiscan ELISA reader.

Three assays were performed for each sample, and the mean reading was calculated.

The results obtained with samples preincubated for 0, 1, 4, 16 and 20 hours are shown in Figure 5. The results demonstrate that the HLA-A2/gag complexes have appreciable stability in culture media at 37°C, with an estimated half-life in excess of 24 hours.

In storage at 0.5-lmg/ml at 4°C HLA-A2/gag complexes appear to be stable for at least 12 months (data not shown).

To demonstrate the ability of the attaching means to cause display of MHC class I on the surface of Daudi cells, Daudi cells deficient in MHC class I were sequentially incubated at 4°C with biotinylated anti-CD20 (Ancell, Nottingham, UK; mAb 2H7 (Berenson et al, Blood 67, 1986, 509-515) at 1ug/ml for 30 minutes); hen egg avidin (S.P.A., Milan, Italy, at 10ug/ml for 10 minutes); biotinylated HLA-A2/gag (at 10ug/ml for 10 minutes); and FITC labelled anti-MHC class I (Ancell, Nottingham, UK; mAB 3F10 (Eisenbarth et al, J Immunol. 124, 1980,1237-1244) at 10 ug/ml). Parallel controls omitted one or other incubation. Cells were washed 3 times in PBS between stages and then fixed in PBS plus 2% formaldehyde and analysed by flow cytometry.

Cells incubated with all three layers of the labelling system had high levels of detectable MHC class I/peptide on their surface compared to untreated Daudi cells (Figure 6). Cells treated with only any 2 components of the 3-step system gave fluorescence levels comparable to untreated cells (data not shown).

5

A chromium release cytotoxicity assay was carried out to establish the ability of specific T cell clones to lyse Daudi or SK-mel-29 cells in accordance with the present invention. Daudi or SK-mel-29 cells were incubated with  $^{51}\text{CrO}_4$  at  $2\mu\text{Ci}/\mu\text{L}$  for 1 hour at  $37^\circ\text{C}$  and then sequentially incubated with: the

10 biotinylated mAbs 2H7 or 225.28s (anti-HMW-MAA) respectively; avidin; and biotinylated HLA-A2/gag complexes as detailed above. Peptide pulsed target cells were incubated with gag 77-85 or melan-A 26-35 peptides at  $0.1\mu\text{M}$  for 1 hour at  $37^\circ\text{C}$ . After washing, labelled target cells were plated into 96-well round bottom plates at 2,500 cells per well, followed by human CTLs at

15 various effector:target ratios. Following incubation at  $37^\circ\text{C}$ , 20ul of supernatant was collected and the amount of  $^{51}\text{Cr}$  released was determined. The percentage of cytotoxicity (lysis) obtained at each effector:target ratio was calculated as:

$100 \times (\text{E}-\text{M})/(\text{T}-\text{M})$ , where E = Experimental release, M = Release in media and T = Release in 5% Triton X-100 detergent.

20

The results shown in Figure 7 are the mean of experiments performed in

duplicate. As shown by these results, the CTL clone (010) efficiently lysed HLA-A2-positive targets (.221/A2) only when these were pre-incubated with the HLA-A2/gag peptide. MHC class I-negative Daudi cells, when targeted with the HLA-A2/gag complexes of the present invention, were recognised and  
5 lysed by this CTL clone to an equivalent degree. Untargeted Daudi cells and cells targeted with only 2 of the 3 components of the targeting system were not recognised (maximal lysis <4% at E:T ratios of up to 80:1).

Control CTL, showing a different HLA-A2-restricted specificity (HLA--  
10 A2/melan-A), did not lyse Daudi cells targeted with the HLA-A2/gag complexes (Fig. 8), demonstrating the fine specificity of the targeting approach.

Untreated Daudi cells pulsed with gag peptide alone were not lysed by clone 010 (data not shown), in keeping with their lack of endogenous MHC class I.

15

The ability of antibody-directed HLA-A2/gag complexes to sensitise the melanoma cell line SK-mel-29 to lysis by HLA-A2/gag-specific CTL line is shown in Figure 8. At all E:T ratios, melanoma cells targeted by complexes linked to surface proteins were lysed substantially more than controls exposed  
20 to only two components of the 3-step targeting system. Additionally, MM9 melanoma cells that do not express HLA-A2 were also lysed in a similar manner (data not shown).

## CLAIMS

1        A complex comprising an HLA class I molecule or fragment thereof,  
which HLA class I molecule or fragment thereof comprises a T cell binding  
5        portion, and attaching means for selectively attaching said HLA class I molecule  
or fragment thereof to a target cell.

2        A complex as claimed in claim 1, wherein said attaching means comprises  
a linking polypeptide with high specific affinity for a target cell specific molecule  
10        on the surface of the target cell.

3        A complex as claimed in claim 2, wherein said linking polypeptide  
comprises an antibody, preferably a monoclonal antibody, raised against said  
target cell specific molecule.

15

4        A complex as claimed in claim 2 or claim 3, wherein said linking  
polypeptide is adapted to be attached directly to said HLA class I molecule or  
fragment thereof.

20        5        A complex as claimed in claim 2 or claim 3, wherein said attaching means  
further comprises a coupling system for coupling said linking polypeptide to said

HLA class I molecule or fragment thereof.

6 A complex as claimed in claim 5, wherein said coupling system comprises  
a two- or three-step chain of well-characterised paired small molecules, which  
5 chain is joined to the linking polypeptide and the HLA class I molecule so as to  
form a stable bridge between the two.

7 A complex as claimed in claim 6, characterised in that said chain  
comprises biotin and avidin/streptavidin.

10

8 A complex as claimed in claim 6, characterised in that said chain  
comprises calmodulin and calmodulin binding peptide.

9 A complex as claimed in any preceding claim, which complex comprises  
15 a recombinant protein, which recombinant protein includes a moiety comprising  
said HLA class I molecule or fragment thereof, and a moiety comprising said  
attaching means.

10 A complex as claimed in any preceding claim, characterised in that said  
20 HLA class I molecule or fragment thereof binds or is attached to a recognition  
peptide, which recognition peptide is arranged to be presented by said HLA class

I molecule or fragment thereof for T cell recognition.

11 A complex as claimed in any preceding claim, characterised in that said  
target cell is a type of cell the presence of which is undesirable in a patient, such  
5 as a tumour cell or a diseased, foreign or malignant cell such as a cancer cell, a  
leukaemia cell, a cell infected with the HIV virus or with any other parasite,  
bacterium, microbe or virus, or a cell responsible for detrimental activity in auto-  
immune disease.

10 12 A complex as claimed in claim 11 appended to claim 10, wherein said  
recognition peptide comprises a peptide which has a strong cytotoxic T cell  
response or which is capable of inducing a powerful immune response.

13 A complex as claimed in claim 11 appended to claim 10, or claim 12,  
15 wherein said recognition peptide comprises a viral or microbial peptide, such as  
an influenza virus peptide, a measles virus peptide, an Epstein-Barr virus peptide,  
in particular an Epstein-Barr virus peptide comprising the RAKFFQLL epitope  
of the lytic protein BZLF1, a Cytomegalovirus peptide, or a tetanus toxoid  
peptide.

20

14 A complex as claimed in any of claims 11-13, wherein the allotype of said



HLA class I molecule or fragment thereof is different from the allotype of the HLA class I molecules of the patient, so that an alloreactive response can additionally or alternatively be triggered against said target cell.

5 15 A complex as claimed in any of claims 1-10, wherein said target cell is an antigen presenting cell.

16 A complex as claimed in claim 15 appended to claim 10, wherein said recognition peptide comprises a tumour specific peptide, or a viral peptide, or a  
10 bacterial peptide, or a parasitic peptide, or any peptide which is exclusively or characteristically presented by HLA class I molecules on the surface of diseased or malignant cells, or virally, bacterially, parasitically or microbially infected cells, or foreign cells the presence of which is undesirable in a patient.

17 A complex as claimed in any preceding claim, wherein said target cell is  
15 a culture cell.

18 A complex as claimed in any of claims 1-16, wherein said target cell is a cell in the body of a patient.

20 19 A method for attaching the complex of any of claims 1-18 to said target cell, comprising the step of introducing to said target cell said HLA class I

molecule or fragment thereof and said attaching means.

20      Use of the complex of claim 15 or claim 16 in the in vivo or ex vivo  
amplification of cytotoxic T cells with specificity for said recognition peptide.

5

21      A method for producing or enhancing an immunological response against  
a target cell, comprising the step of attaching the complex of any of claims 1-18  
to said target cell in accordance with the method of claim 19.

10      22      A method for immunising a patient against a disease or condition which  
is characterised by the presence in the patient's body of cells displaying said  
recognition peptide on the surface thereof; such as a tumour, or a malignant or  
auto-immune disease such as cancer or leukaemia, an infectious disease such as  
a viral infection such as HIV infection, a bacterial or microbial infection such as  
15      tuberculosis, or a parasitic infection such as malaria; comprising the step of  
administering to said patient an effective amount of the complex of claim 15 or  
claim 16.

23      A pharmaceutical composition for use in immunising a patient against a  
20      disease or condition which is characterised by the presence in the body of the  
patient of diseased, malignant or foreign cells; such as a tumour, or a malignant

or auto-immune disease such as cancer or leukaemia, or an infectious disease such as a viral infection such as HIV infection, or a bacterial or microbial infection such as tuberculosis, or a parasitic infection such as malaria; said pharmaceutical composition comprising a complex as claimed in claim 15 or claim 16 and an appropriate excipient or carrier.

24 Use of the complex of claim 15 or claim 16 in the preparation of a medicament for use in immunising a patient against a disease or condition which is characterised by the presence in the patient's body of cells displaying said recognition peptide on the surface thereof; such as a tumour, or a malignant or auto-immune disease such as cancer or leukaemia, an infectious disease such as a viral infection such as HIV infection, a bacterial or microbial infection such as tuberculosis, or a parasitic infection such as malaria.

15 25 A method for the treatment of a disease or condition such as a tumour, or a malignant or auto-immune disease such as cancer or leukaemia, an infectious disease such as a viral infection such as HIV infection, a bacterial or microbial infection such as tuberculosis, or a parasitic infection such as malaria, comprising the step of administering to a patient in need thereof an effective amount of the complex of any of claims 11-14.

26 A pharmaceutical composition for use in the treatment of a disease or condition characterised by the presence in a patient of diseased, foreign or malignant cells; such as a tumour, or a malignant or auto-immune disease such as cancer or leukaemia, or an infectious disease such as a viral infection such as HIV infection, or a bacterial or microbial infection such as tuberculosis, or a parasitic infection such as malaria; said pharmaceutical composition comprising a complex as claimed in any of claims 11-14 and an appropriate excipient or carrier.

10 27 Use of the complex of any of claims 11-14 in the preparation of a medicament for the treatment of a tumour, or a malignant or auto-immune disease such as cancer or leukaemia, or an infectious disease such as a viral infection such as HIV infection, or a bacterial or microbial infection such as tuberculosis, or a parasitic infection such as malaria.

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28 A pharmaceutical pack or kit comprising one or more containers containing one or more of the pharmaceutical compositions claimed in claim 23 or claim 26 and written instructions for the administration of said composition or compositions to a patient.

20



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Claims searched: 1-19, 21, 23, 24, 26-28;      Date of search: 25 November 1999  
20 (in part)

**Patents Act 1977**  
**Search Report under Section 17**

**Databases searched:**

UK Patent Office collections, including GB, EP, WO & US patent specifications, in:

UK Cl (Ed.Q):

Int Cl (Ed.6):

Other: ONLINE: WPI, EPODOC, PAJ, BIOSIS, CAPLUS, BIOTECHABS,  
CANCERLIT, EMBASE, LIFESCI, MEDLINE

**Documents considered to be relevant:**

Category	Identity of document and relevant passage	Relevant to claims
X	EP 0352761 A2 (BEHRINGWERKE) Whole document	1, at least
X	WO 97/35991 A1 (JOHNS HOPKINS UNI.) page 9 lines 7-10; page 17 line 28-page 18 line 13; page 33 line 27-page 34 line 11; Fig.1C	1, at least
X	WO 97/28191 A1 (DADE INT.) page 49 line 17-page 55 line 8; page 108 lines 11-14; Fig. 1C	1, at least
X	WO 97/24446 A2 (CHIRON VIAGENE INC.) page 2 lines 10-13; page 3 lines 7-10; page 9 lines 13-24; Examples 2-4; Figs. 2, 5 & 9	1, at least
X	WO 93/10220 A1 (ANERGEN INC.) page 3 lines 6-14; page 4 lines 28-30	1, at least

X	Document indicating lack of novelty or inventive step	A	Document indicating technological background and/or state of the art.
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